The Fisher & Paykel Healthcare classic circuit can be connected to the Neopuff™ or other T-Piece resuscitator.* It features an adjustable PEEP valve, and can be connected to a resuscitation mask or endotracheal tube.

*If the T-Piece resuscitator meets the gas-powered resuscitator standard (ISO 10651-5:2006)
PRODUCT SPECIFICATIONS

Compatible with:

T-Piece resuscitators: Neopuff® RD900 series or other T-Piece resuscitator that meets the gas-powered resuscitator standard (ISO 10651-5:2006)

F&P resuscitation masks: RD803 (XS); RD804 (S); RD805 (M); RD806 (L); RD807 (XL)

Connection: 15 mm medical taper at patient connection; 10 mm medical taper at Neopuff connection

Purchasing options (Boxes of 10):

<table>
<thead>
<tr>
<th>Product code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>900RD010</td>
<td>Classic T-Piece circuit</td>
</tr>
<tr>
<td>900RD014-10</td>
<td>Classic T-Piece circuit + 42mm diameter mask (S)</td>
</tr>
<tr>
<td>900RD015-10</td>
<td>Classic T-Piece circuit + 50mm diameter mask (M)</td>
</tr>
<tr>
<td>900RD016-10</td>
<td>Classic T-Piece circuit + 60mm diameter mask (L)</td>
</tr>
</tbody>
</table>

PERFORMANCE

Resistance to flow (average): 0.6 cmH₂O at 15 L/min

Min flow rate (L/min): 5 L/min

Max flow rate (L/min): 15 L/min

Length of tubing: 1.6 m

Internal diameter: Nominal 12 mm

Compliance (average): 2.23 ml/kPa/m

Ambient range °C: -18 °C to 50 °C

Duration of use: Single patient use – supplied clean, not sterile

Positive End Expiratory Pressure (PEEP)*:

@ 5 L/min: 1 to 6 cmH₂O [mbar]

@ 8 L/min: 1 to 10 cmH₂O [mbar]

@ 10 L/min: 2 to 15 cmH₂O [mbar]

@ 15 L/min: 4 to 17 cmH₂O [mbar]

* All performance figures listed above are representative only.
PEEP values stated are based on typical clinical PIP settings. Higher PEEP values can be achieved if higher PIP values are set.

COMPONENTS AND COMPOSITIONS

Predominant materials: Polycarbonate, acetal, polystyrene, polyethylene, stainless steel

Materials not present: Not manufactured with natural rubber latex, PVC or Phthalates (DEHP, DBP, BBP)

Manufacturing mode: Produced in a controlled working environment

Disposal: According to hospital protocol

Shelf life: 3 years

REGULATORY

Classification: Class IIa (EU and Australia), Class II (Canada), Class I (USA)

Country of origin: New Zealand

Notified body: TÜV SÜD Product Service GmbH, 0123

PEEP: positive end-expiratory pressure; PIP: peak inspiratory pressure

Please note that the information in this specifications catalogue (including product information and images) is summarized and provided for illustrative purposes only. Please refer to the relevant user instructions for more information and confirm details with your local Fisher & Paykel Healthcare representative prior to placing an order. Information subject to change without notice.

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